

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
22 May 2003 (22.05.2003)

PCT

(10) International Publication Number
WO 03/042283 A1

(51) International Patent Classification⁷: **C08G 77/42**,
77/392, C08B 15/05, 33/00, 37/00, C07F 7/08

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(21) International Application Number: PCT/JP02/11807

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(22) International Filing Date:

12 November 2002 (12.11.2002)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

NO.2001-348122

13 November 2001 (13.11.2001) JP

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

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Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.



WO 03/042283 A1

(54) Title: SACCHARIDE RESIDUE-FUNCTIONAL ORGANOPOLYSILOXANES AND METHOD FOR THE PREPARATION THEREOF

(57) Abstract: A saccharide residue-functional organopolysiloxane containing at least two monosaccharide or polysaccharide groups per molecule in which a specific site on the monosaccharide or polysaccharide is bonded to silicon through a thioether bond. Also, a method of preparing the saccharide residue-functional organopolysiloxane, comprising condensing a saccharide residue-functional metal thiolate compound wherein the metal is an alkali metal atom or alkaline-earth metal atom, and an organopolysiloxane containing groups having the formula -R²Q wherein R² is C₂ to C₁₀ alkylene, and Q is a group selected from halogen atoms, C₁ to C₁₀ alkylsulfonate groups, and C₆ to C₂₀ arylsulfonate groups.

DESCRIPTION

SACCHARIDE RESIDUE-FUNCTIONAL ORGANOPOLYSILOXANES AND
METHOD FOR THE PREPARATION THEREOF

5

[0001] This invention relates to novel saccharide residue-functional organopolysiloxanes and to a method for their preparation. More particularly, this invention relates to saccharide residue-functional organopolysiloxanes in which monosaccharide or polysaccharide is bonded through a thioether bond to silicon and to a method for the preparation of said
10 saccharide residue-functional organopolysiloxanes.

[0002] While the important role played by saccharides in mediating biological functions has been known for some time, unique interactions between substances originating in the steric structure of saccharides have been elucidated only very recently, and, concurrent therewith, the development of drugs and functional materials that effectively utilize these
15 functionalities has attracted attention.

[0003] With this goal in mind, methods for bonding saccharides to synthetic polymers are already known, as are attempts to apply these methods. Organopolysiloxanes, because they are biologically inert, are ideal materials for drugs, therapeutic materials, and cosmetics, and a number of saccharide residue-functional organopolysiloxanes — and methods for their
20 preparation — have been introduced to date.

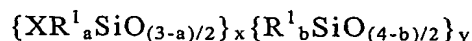
[0004] For example, Japanese Patent Application Publication (Kokai) No. Sho 62-68820 (68,820/1987) discloses saccharide residue-functional organopolysiloxane in which saccharide is linked to polysiloxane through an amide bond; Japanese Patent Application Publication (Kokai) No. Hei 5-186596 (186,596/1993) discloses saccharide residue-
25 functional organopolysiloxane in which saccharide is linked to polysiloxane through a glycosidic bond; Japanese Patent Application Publication (Kokai) No. Hei 8-134103 (134,103/1996) discloses saccharide residue-functional organopolysiloxane in which saccharide is linked to polysiloxane through a urethane bond; and Japanese Patent Application Publication (Kokai) No. Hei 11-92490 (92,490/1999) discloses saccharide
30 residue-functional organopolysiloxane in which saccharide is linked to polysiloxane through a glycosidic or thioglycosidic bond.

[0005] These saccharide residue-functional organopolysiloxanes, however, suffer from a number of drawbacks. For example, due to instability in the linkage moiety, cleavage of the

linkage resulting in decomposition can occur quite readily in these saccharide residue-functional organopolysiloxanes. In addition, depending on the particular linkage moiety, the polar interactions of the linkage moiety can inhibit the manifestation of the properties of the saccharide. With regard to the synthesis of the referenced saccharide residue-functional organopolysiloxanes, it has been quite difficult to effect bonding at only a specific site of the saccharide.

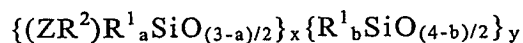
[0006] The inventors achieved this invention as a result of intensive investigations directed to solving the problems identified above. In specific terms, an object of this invention is to provide novel saccharide residue-functional organopolysiloxanes that are stable and that do not suffer from any impairment in the functionality of the saccharide. An additional object of this invention is to provide a method for the preparation of these novel saccharide residue-functional organopolysiloxanes. More particular objects of this invention are to provide novel saccharide residue-functional organopolysiloxanes in which a specific site on the monosaccharide or polysaccharide is bonded through a thioether bond to silicon and to provide a method for the preparation of these novel saccharide residue-functional organopolysiloxanes.

[0007] This invention relates to saccharide residue-functional organopolysiloxane that is represented by the general formula:



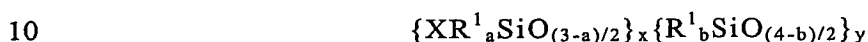
wherein R^1 is C_1 to C_{10} alkyl or aryl; X is a group having the formula $-R^2-S-R^3-Y$ wherein R^2 and R^3 are each independently selected from C_1 to C_{20} divalent hydrocarbyl and Y is a substituted or unsubstituted monosaccharide or polysaccharide residue having an oxygen atom bonded to R^3 ; a is an integer from 0 to 2; b is an integer from 0 to 3; x is an integer with a value of at least 2; and y is an integer with a value of at least 0; wherein when an individual siloxane structural unit is present in a plural number, these may be the same as or may differ from each other.

[0008] This invention additionally relates to a method for preparing the saccharide residue-functional organopolysiloxane, by effecting a condensation reaction between a saccharide residue-functional metal thiolate compound having the formula $M-S-R^3-Y$ wherein R^3 and Y are defined as above, M is an alkali metal atom or alkaline-earth metal atom, and an organopolysiloxane represented by the general formula:



wherein R^1 , R^2 , a , b , x , and y are defined as above and Z is a group selected from halogen atoms, C_1 to C_{10} alkylsulfonate groups, and C_6 to C_{20} arylsulfonate groups wherein when an individual siloxane structural unit is present in a plural number, these may be the same as or may be different from each other.

[0009] The saccharide residue-functional organopolysiloxane according to this invention is represented by the following general formula:



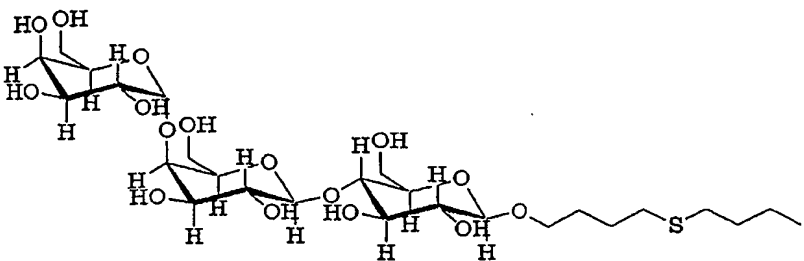
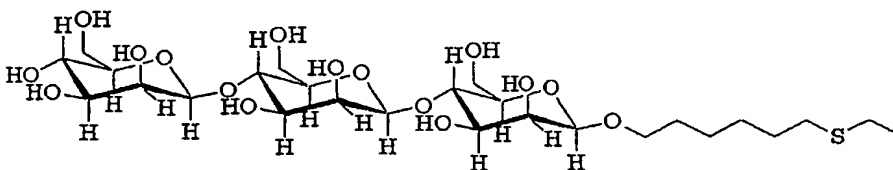
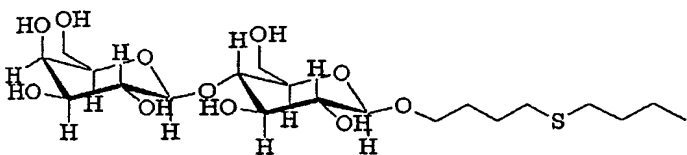
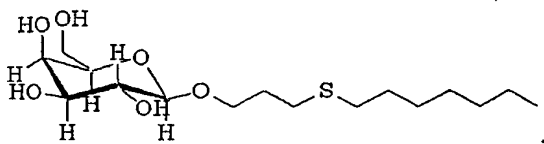
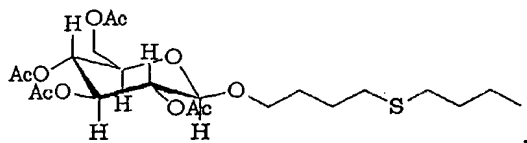
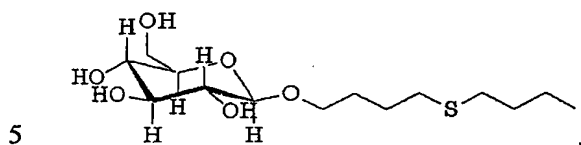
R^1 in this formula is C_1 to C_{10} alkyl or aryl. The alkyl can be exemplified by methyl, ethyl, propyl, butyl, pentyl, isopropyl, isobutyl, cyclopentyl, and cyclohexyl, while the aryl can be exemplified by phenyl and naphthyl. Methyl is preferred among the preceding for R^1 . X in the preceding formula is a group with the general formula $-R^2-S-R^3-Y$ and is bonded to the silicon atom in the organosiloxane. R^2 and R^3 in this group are each independently selected from C_1 to C_{20} divalent hydrocarbonyl and can be exemplified by straight-chain alkylene such as ethylene, propylene, butylene, and hexylene; branched alkylene such as methylethylene, methylethylene, 1-methylpentylene, and 1,4-dimethylbutylene; arylene such as 1,4-phenylene, 1,3-phenylene, and 4,4'-biphenylene; and aralkylene such as 1,4-bis(methylene)phenylene and 2,2-(para-phenylene)propylene. Y in $-R^2-S-R^3-Y$ represents a substituted or unsubstituted, mono- or polysaccharide residue whose bonding site with R^3 is in all instances an oxygen atom.

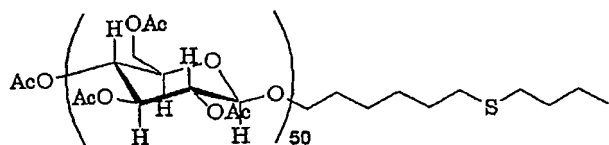
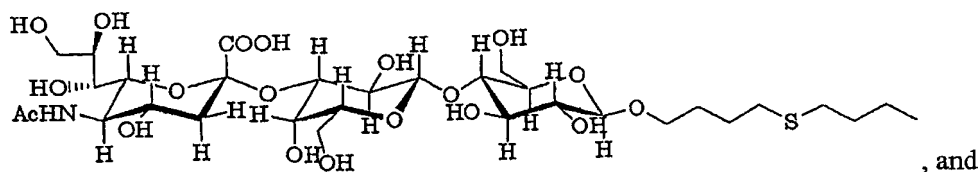
[0010] Monosaccharide units that can form the saccharide structure can be exemplified by glucopyranose (glucose), mannose, allose, altrose, galactose, idose, talose, gulose, ribose, arabinose, xylose, fructose, fucose, N-acetylglucosamine, N-acetylgalactosamine, sialic acid, and esters of the preceding. Y can be specifically exemplified by monosaccharide residues such as glucopyranosyl (glucosyl), mannosyl, allosyl, altrosyl, galactosyl, idosyl, talosyl, gulosyl, ribosyl, arabinosyl, xylosyl, fucosyl, fructosyl, N-acetylglucosaminyl, N-acetylgalactosaminyl, sialyl, and their esters; oligosaccharide residues such as mantosyl, cellobiosyl, lactosyl, mannotriosyl, globotriosyl, and their esters; and polysaccharide residues such as cellulosyl and amylosyl and their esters. Preferred among the preceding are monosaccharide residues and oligosaccharide residues that contain no more than 5

4

saccharide units. In addition, the oxygen atom in Y that bonds with R³ is preferably a glycosidic oxygen atom.

[0011] The X under consideration can be specifically exemplified by structures having the following formulas wherein Ac is acetyl:





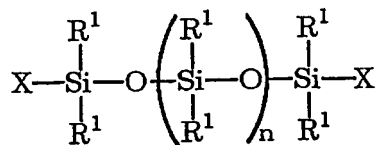
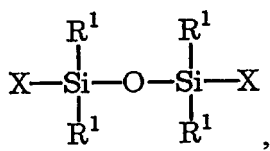
5 [0012] The subscripts in the general formula:

$$\{XR^1_aSiO_{(3-a)/2}\}_x\{R^1_bSiO_{(4-b)/2}\}_y$$

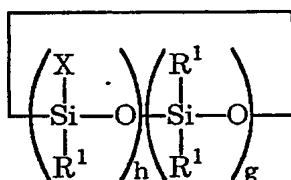
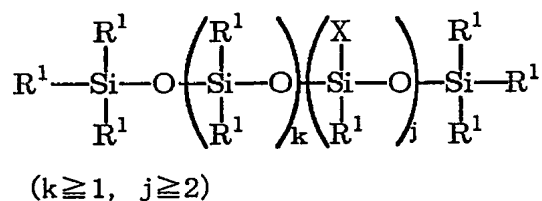
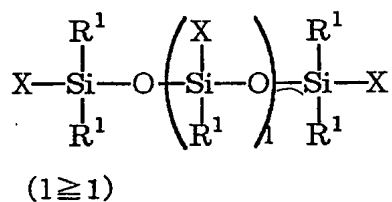
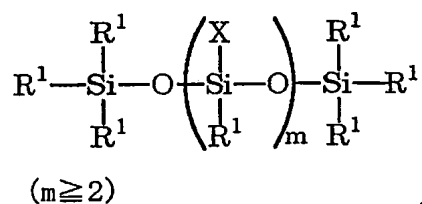
10 for the inventive saccharide residue-functional organopolysiloxane have the following values: a is an integer from 0 to 2, b is an integer from 0 to 3, x is an integer with a value of at least 2, and y is an integer with a value of at least 0. When an individual siloxane structural unit is present in a plural number, these may be the same as or may differ from each other. The organopolysiloxane under consideration can be specifically exemplified by organopolysiloxanes with the following general formulas, in which X and R¹ are defined as

15 above and e, f, g, h, j, k, l, m, and n are integers that indicate the number of siloxane units present in each molecule:

[0013]



20 (n ≥ 1)



($h \geq 2, g \geq 1$)

$\text{R}^1_f\text{Si}(\text{OSiR}^1_2\text{X})_{4-f}$ ($f = 0, 1$),

$\text{X}_e\text{Si}(\text{OSiR}^1_2\text{X})_{4-e}$ ($e = 0, 1$),

$(\text{XSiO}_{3/2})_8$, and

10 $(\text{SiO}_{4/2})_8(\text{XR}^1_2\text{SiO}_{1/2})_8$.

[0014] The inventive saccharide residue-functional organopolysiloxane can be synthesized by effecting a condensation reaction between a saccharide residue-functional metal thiolate compound $\text{M}-\text{S}-\text{R}^3-\text{Y}$ and organopolysiloxane represented by the general formula

$\{(\text{ZR}^2)\text{R}^1_a\text{SiO}_{(3-a)/2}\}_x\{(\text{R}^1_b\text{SiO}_{(4-b)/2})_y$, wherein $\text{R}^1, \text{R}^2, \text{R}^3, \text{Y}, a, b, x$, and y are defined as above and ZR^2 is bonded to the silicon atom in the organosiloxane. Z is a group selected from halogen atoms, C_1 to C_{10} alkylsulfonate groups, and C_6 to C_{20} arylsulfonate groups.

The halogen atoms encompassed by Z can be exemplified by the chlorine atom, bromine

atom, and iodine atom. The alkylsulfonate group can be exemplified by methanesulfonate and ethanesulfonate and the arylsulfonate group can be exemplified by benzenesulfonate and toluenesulfonate. M is an alkali metal or alkaline-earth metal atom, for example, lithium, sodium, potassium, calcium, or magnesium. Since saccharide residue-functional metal thiolate compounds $M-S-R^3-Y$ are generally difficult to isolate, they are preferably synthesized in the reaction system, for example, by the action of a metal or metal hydride on the corresponding saccharide residue-functional thiol compound or by the action of a metal on the corresponding saccharide residue-functional thioester or thioether compound, and used directly for the condensation reaction.

- 10 [0015] The method for preparing the organopolysiloxane starting material $\{(ZR^2)R^1_aSiO_{(3-a)/2}\}_x\{R^1_bSiO_{(4-b)/2}\}_y$ used in the synthesis of the inventive saccharide residue-functional organopolysiloxane is not critical. For example, this starting material can be directly synthesized by a hydrosilylation reaction between a halogen-, alkylsulfonate-, or arylsulfonate-functional alkenyl compound and a known SiH-functional
- 15 organopolysiloxane. Alternatively, utilizing the hydroxyl present in an organopolysiloxane, this starting material can be synthesized by the introduction of halogen by a substitution reaction or by the introduction of an alkylsulfonate or arylsulfonate group by an esterification reaction. The starting material under consideration can also be synthesized by preparing the organopolysiloxane by the hydrolysis of an organohalosilane or
- 20 organoalkoxysilane bearing a halogen-, alkylsulfonate-, or arylsulfonate-substituted organic group.

- [0016] The saccharide residue-functional metal thiolate compound $M-S-R^3-Y$ can be synthesized, for example, by converting the glycosidic hydroxyl of the saccharide molecule to an alkenyl ether by a known method; addition reacting a thiocarboxylic acid or thiol
- 25 compound with the alkenyl group in the presence of a radical initiator to give, respectively, a thioester derivative or thioether compound; and reacting these products, respectively, with an alkali metal base such as alkali metal hydroxide or alkali metal methoxide, with alkaline-earth metal base such as alkaline-earth metal hydroxide, with an alkali metal, or with alkaline-earth metal.

- 30 [0017] The inventive saccharide residue-functional organopolysiloxane as described hereinabove, because it characteristically has a structure in which a specific site in the monosaccharide or polysaccharide is bonded to silicon through a thioether bond, is therefore also characterized by its stable nature and by no loss of saccharide functionality. As a

consequence, the inventive saccharide residue-functional organopolysiloxane exhibits an excellent durability as a material while at the same time retaining the properties inherent to saccharides, such as pharmacological activity and biocompatibility. The inventive saccharide residue-functional organopolysiloxane therefore offers the advantage of being useful in a variety of applications, for example, as an ingredient for cosmetics, as a reagent for the separation of optical isomers, as a therapeutic material for the separation of, for example, toxins or viruses, as a therapeutic agent, and as an agrochemical. The inventive method for synthesizing saccharide residue-functional organopolysiloxane is characterized by its ability to very efficiently produce the subject organopolysiloxane.

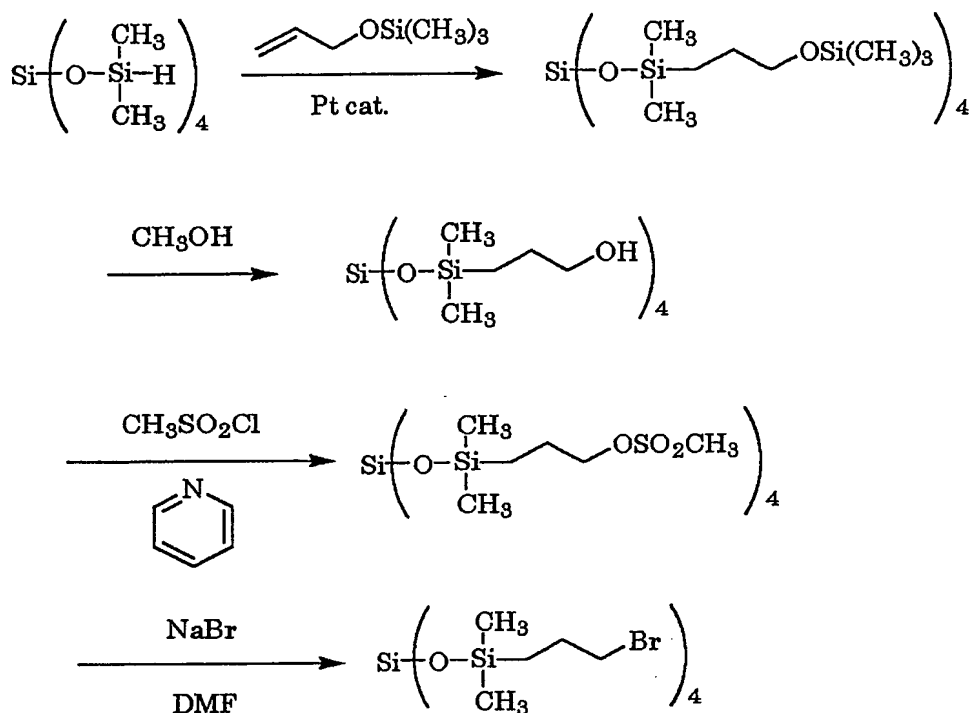
[0018] The invention is explained in greater detail hereinbelow through working examples. The inventive saccharide residue-functional organopolysiloxane was identified in the examples by nuclear magnetic resonance analysis. In the reaction equations, chemical formulas, and text that follow, Pt cat. denotes a complex of platinum and 1,1,3,3-tetramethyl-1,3-divinyldisiloxane, Ac stands for the acetyl group, DMF stands for N,N-dimethylformamide, AIBN stands for azobisisobutyronitrile, and Ph stands for the phenyl group.

[0019]

Reference Example 1

Tetrakis(bromopropyltrimethylsiloxy)silane was synthesized by the following reactions starting from tetrakis(dimethylsiloxy)silane. Thus, allyloxytrimethylsilane was reacted with tetrakis(dimethylsiloxy)silane in the presence of a platinum 1,1,3,3-tetramethyl-1,3-divinyldisiloxane complex as catalyst to produce tetrakis(trimethylsiloxypropyltrimethylsiloxy)silane, from which the trimethylsiloxy group was subsequently eliminated by the action of excess methanol to give tetrakis(hydroxypropyltrimethylsiloxy)silane. Methanesulfonyl chloride was then reacted in pyridine with the tetrakis(hydroxypropyltrimethylsiloxy)silane to give tetrakis(methanesulfonyloxypropyltrimethylsiloxy)silane, which was subsequently reacted with sodium bromide in DMF to give tetrakis(bromopropyltrimethylsiloxy)silane:

[0020]

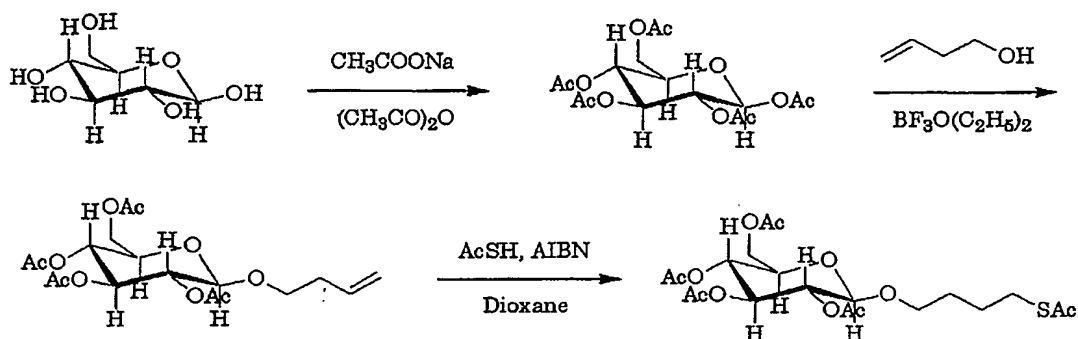


[0021]

Reference Example 2

4-acetylthiobutyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was synthesized by the following reactions from β-D-glucopyranose. Thus, acetyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was prepared by exhaustive acetylation of the hydroxyl groups in β-D-glucopyranose by the action of sodium acetate in acetic anhydride. The acetyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was subsequently reacted with 3-buten-1-ol under catalysis by boron trifluoride diethyl etherate to give butenyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside. The butenyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was mixed with thioacetic acid in dioxane and AIBN was added as radical initiator; reaction then gave 4-acetylthiobutyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside:

[0022]

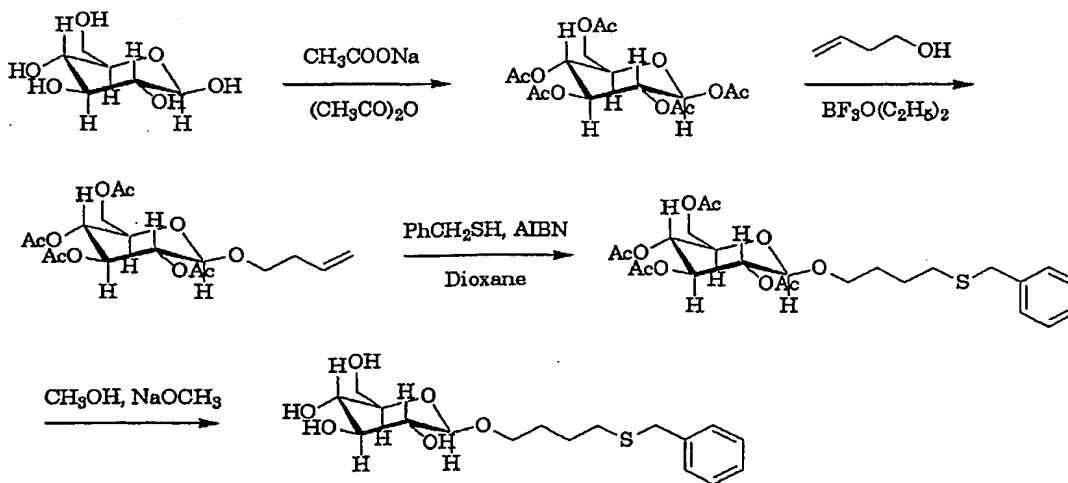


[0023]

Reference Example 3

4-benzylthiobutyl β-D-glucopyranoside was synthesized by the following reactions from β-D-glucopyranose. Thus, acetyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was prepared by exhaustive acetylation of the hydroxyl groups in β-D-glucopyranose by the action of sodium acetate in acetic anhydride. The acetyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was subsequently reacted with 3-buten-1-ol under catalysis by boron trifluoride diethyl etherate to give butenyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside. The butenyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside was mixed with benzyl thiol in dioxane and AIBN was added as radical initiator; reaction then gave 4-benzylthiobutyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside. Deacetylation of the 4-benzylthiobutyl 2,3,4,6-tetra-O-acetyl-β-D-glucopyranoside by reaction with sodium methoxide in methanol gave 4-benzylthiobutyl β-D-glucopyranoside:

15 [0024]

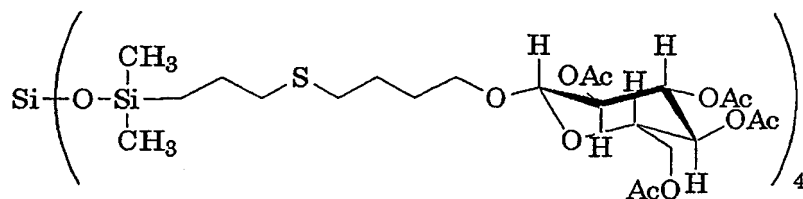


[0025]

Example 1

The tetrakis(bromopropyldimethylsiloxy)silane synthesized in Reference Example 1 (70 mg, 0.086 mmol) and 495 mg (1.034 mmol) of the 4-acetylthiobutyl 2,3,4,6-tetra-O-acetyl-
 5 β -D-glucopyranoside synthesized in Reference Example 2 were dissolved in 0.5 mL anhydrous DMF and 0.5 mL anhydrous methanol, and the liquid mixture was stirred for 2 hours at room temperature. Sodium methoxide (62 mg, 1.137 mmol) was then added and stirring was carried out for another 16 hours at 35 °C. Acetic acid (0.3 mL) was added, stirring was carried out for 10 minutes at room temperature, and 5 mL acetic anhydride and
 10 5 mL pyridine were then added. The reaction solution was subsequently poured into ice water, and this was extracted 3 times with chloroform. The combined organic layers were washed with 1 N hydrochloric acid, saturated aqueous sodium bicarbonate, saturated aqueous sodium chloride, and then dried over anhydrous magnesium sulfate. After concentration, the product was purified by column chromatography and gel permeation
 15 chromatography to give 70 mg of a saccharide residue-functional organopolysiloxane with the following formula:

[0026]



¹H-nuclear magnetic resonance spectral data (solvent: deuteriochloroform) δ = 0.07 (s, 24H), 0.63 (t, J = 9 Hz, 8H), 1.5-1.7 (m, 24H), 1.99 (s, 12H), 2.00 (s, 12H), 2.03 (s, 12H), 2.07(s, 12H), 2.4-2.6 (m, 16H), 3.5 (m, 4H), 3.7 (m, 4H), 3.9 (m, 4H), 4.1 (m, 4H), 4.3 (m, 4H), 4.48 (d, J = 8 Hz, 4H), 4.97 (t, J = 10 Hz, 4H), 5.07 (t, J = 10 Hz, 4H), 5.17 (t, J = 10 Hz, 4H).

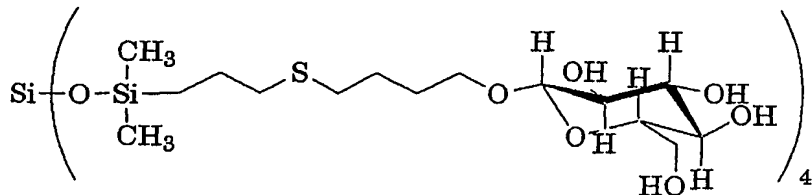
[0027]

25 Example 2

The product obtained in Example 1 (55 mg, 0.025 mmol) was dissolved in 1 mL dry methanol, 0.53 mg (0.0098 mmol) sodium methoxide was added, and stirring was carried out for 24 hours at room temperature. Cation-exchange resin (0.2 mL, Amberlite IR-120B H AG) was added to the resulting reaction mixture followed by stirring and then filtration.

Concentration of the filtrate in vacuo gave 38 mg of the a saccharide residue-functional organopolysiloxane having the following formula:

[0028]



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¹H-nuclear magnetic resonance spectral data (solvent: deuterium oxide) δ = 0.0 (s, 24H), 0.57 (t, J = 9 Hz, 8H), 1.5-1.7 (m, 24H), 2.4-2.5 (m, 16H), 3.1 (m, 4H), 3.2-3.4 (m, 12H), 3.5-3.7 (m, 8H), 3.7-3.9 (m, 8H), 4.30 (d, J = 8 Hz, 4H).

[0029]

10 Example 3

The 4-benzylthiobutyl β -D-glucopyranoside synthesized in Reference Example 3 (353 mg, 0.985 mmol) was cooled to -35 °C, ammonia gas was bubbled in so as to liquefy 30 mL ammonia, 226 mg (9.85 mmol) sodium metal was added, and stirring was carried out for 30 minutes at -35 °C. This was followed by the addition of 474 mg (8.86 mmol)

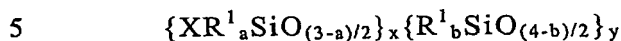
15 ammonium chloride and 100 mg (0.123 mmol) of the tetrakis(bromopropyldimethylsiloxy)silane synthesized in Reference Example 1 dissolved in 3 mL dimethoxyethane, and while stirring the temperature was then gradually returned to room temperature with evaporation of ammonia gas. After concentration, the product was purified by column chromatography and gel permeation chromatography to give 21 mg of a

20 saccharide residue-functional organopolysiloxane having the same structure as the product in Example 2.

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CLAIMS

1. A saccharide residue-functional organopolysiloxane having the formula:



wherein R^1 is C_1 to C_{10} alkyl or aryl; a is an integer from 0 to 2; b is an integer from 0 to 3; x is an integer with a value of at least 2; y is an integer with a value of at least 0; and X is a group having the formula $-R^2-S-R^3-Y$ wherein R^2 and R^3 are each independently selected from C_1 to C_{20} divalent hydrocarbyl and Y is a substituted or unsubstituted monosachharide or polysaccharide residue having an oxygen atom bonded to R^3 .

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2. The saccharide residue-functional organopolysiloxane according to claim 1, wherein R^1 is methyl.

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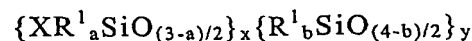
3. The saccharide residue-functional organopolysiloxane according to claim 1, wherein Y is a monosaccharide residue.

4. The saccharide residue-functional organopolysiloxane according to claim 1, wherein Y is an oligosaccharide residue containing no more than five saccharide units.

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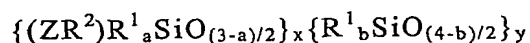
5. The saccharide residue-functional organopolysiloxane according to claim 1, wherein the oxygen atom in Y bonded to R^3 is a glycosidic oxygen atom.

25 6. A method of preparing a saccharide residue-functional organopolysiloxane having the formula:



30 wherein R^1 is C_1 to C_{10} alkyl or aryl; a is an integer from 0 to 2; b is an integer from 0 to 3; x is an integer with a value of at least 2; y is an integer with a value of at least 0; and X is a group having the formula $-R^2-S-R^3-Y$ wherein R^2 and R^3 are each independently selected from C_1 to C_{20} divalent hydrocarbyl and Y is a substituted or unsubstituted monosachharide

or polysaccharide residue having an oxygen atom bonded to R^3 , comprising condensing a saccharide residue-functional metal thiolate compound having the formula $M-S-R^3-Y$ wherein R^3 is C_1 to C_{20} divalent hydrocarbyl, Y is a substituted or unsubstituted monosachharide or polysaccharide residue having an oxygen atom bonded to R^3 , and M is an alkali metal atom or alkaline-earth metal atom, and an organopolysiloxane having the formula:



wherein R^1 is C_1 to C_{10} alkyl or aryl; R^2 is C_1 to C_{20} divalent hydrocarbyl; a is an integer from 0 to 2; b is an integer from 0 to 3; x is an integer with a value of at least 2; y is an integer with a value of at least 0; and Z is a group selected from halogen atoms, C_1 to C_{10} alkylsulfonate groups, and C_6 to C_{20} arylsulfonate groups.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/JP 02/11807

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C08G77/42 C08G77/392 C08B15/05 C08B33/00 C08B37/00
C07F7/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C08G C08B C07F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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X	PATENT ABSTRACTS OF JAPAN vol. 013, no. 485 (M-887), 6 November 1989 (1989-11-06) & JP 01 192490 A (MAZDA MOTOR CORP), 2 August 1989 (1989-08-02) cited in the application abstract ----- -/--	1

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
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- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

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- * & * document member of the same patent family

Date of the actual completion of the international search

17 February 2003

Date of mailing of the international search report

05/03/2003

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	WO 00 08038 A (SYNSORB BIOTECH INC ;HINDSGAUL OLE (CA); HUMMEL GERD (DE)) 17 February 2000 (2000-02-17) claims 1,5 page 14, line 18 - line 26 ----	1
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